Amendments to the Claims

Please cancel claims 5-10, 13 and 14 without prejudice. Please amend claims 1, 4, 11, 12, 15, and 16 and add new claims 17 and 18 as set forth below in the List of Claims.

List of Claims

1. (Currently amended A compound of the formula I:

OH
$$H_2N$$
 NH R^6 R^8 R^8 R^9 NH_2 R^3 CH_2 CH_2 CH_2 CH_2 CH_2 CH_3 CH_4 CH_5 CH_5 CH_6 CH_7 CH_8 CH_8

wherein

R¹ is selected from

- (i) <u>a</u> linear or branched C₁-C₆ alkyl;
- (ii) $\underline{a} C_1 C_6 \text{ alkoxy};$

R² is selected from

- (i) hydrogen;
- (ii) <u>a</u> linear or branched C₁-C₆ alkyl;
- (iii) $\underline{a} C_1 C_6 \text{ alkoxy};$

R³ and R⁴ is each and independently selected from

- (i) hydrogen;
- (ii) \underline{a} linear or branched C_1 - C_6 alkyl;

(iii)
$$-\xi$$
-(CH₂)_m wherein m = 1-3;

$$(iv)$$
 $-\xi$ - (CH_2) ; and

R⁵, R⁶, R⁷, R⁸ and R⁹ is each and independently selected from

- (i) hydrogen;
- (ii) <u>a halogen, where "halogen" encompasses wherein said halogen is selected</u> from the group consisting of: chloro, fluoro, bromo and iodo; and
- (iii) <u>a</u> linear or branched C₁-C₆ alkyl; and

n is an integer of from 1 to 5;

as well as pharmaceutically and pharmacologically acceptable salts thereof.

2. (Original) A compound of formula I according to claim 1, wherein

R¹ is a linear C₁-C₆ alkyl;

R² is a linear C₁-C₆ alkyl or hydrogen;

 R^3 and R^4 is each and independently selected from a straight C_1 - C_6 alkyl or hydrogen; R^5 , R^6 , R^7 , R^8 and R^9 is each and independently selected from

- (i) hydrogen;
- (ii) halogen, where "halogen" encompasses chloro, fluoro, bromo and iodo;
- (iii) linear or branched C₁-C₆ alkyl; and

n is an integer of from 1 to 5.

3. (Original) A compound according to claim 1, wherein

R¹ is CH₃;

R² is hydrogen or CH₃;

R³ and R⁴ are both hydrogen; and

R⁵, R⁶, R⁷, R⁸ and R⁹ are all hydrogen; and

n is 4.

4. (Currently amended) A compound according to claim 1, which wherein said compound is selected from anyone the group consisting of:

H-Dmt-D-Arg-Phe-Lys-NH₂;

H-Dmt-D-Arg-Phe-Orn-NH₂;

H-Dmt-D-Arg-Phe-A₂Bu-NH₂;

H-Mmt-D-Arg-Phe-Lys-NH₂;

H-Dmt-D-Arg-Phe(p-F)-Lys-NH₂; and

Dmt(NMe)-D-Arg-Phe-Lys-NH₂.

5-10. Cancelled

- 11. (Currently amended) A pharmaceutical composition comprising a compound of the formula I according to claim 1 as an active ingredient, in admixture with one or more pharmacologically and pharmaceutically acceptable carriers.
- 12. (Currently amended) A <u>The</u> pharmaceutical composition according to <u>of</u> claim 11, <u>wherein said pharmaceutical composition is</u> suitable for administration intrathecally, epidurally, intramuscularly, and intravenously.
- 13. Cancelled
- 14. Cancelled
- 15. (Currently amended) A method for the treatment of a subject of treating a patient suffering from pain, whereby an effective amount of comprising administering to said patient a compound of the formula I of claim 1, is administered to a patient in need of pain relief according to claim 1 for a time and under conditions effective to induce analgesia.

- 16. (Currently amended) A <u>The</u> method for <u>of</u> treatment according to <u>of</u> claim 15, wherein the <u>said</u> pain is labor pain.
- 17. (New) A salt of a compound according to claim 1 selected from the group consisting of: a hydrochloride, an acetate, or a trifluoroacetate salt.
- 18. (New) A process for preparing a compound of formula I according to claim 1, comprising:
 - a) preparing a peptide attached to a solid phase support;
 - b) coupling a protected amino acid to said peptide in an inert solvent using a coupling agent;
 - c) completing the synthesis; and
 - d) isolating the compound of formula I.